

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



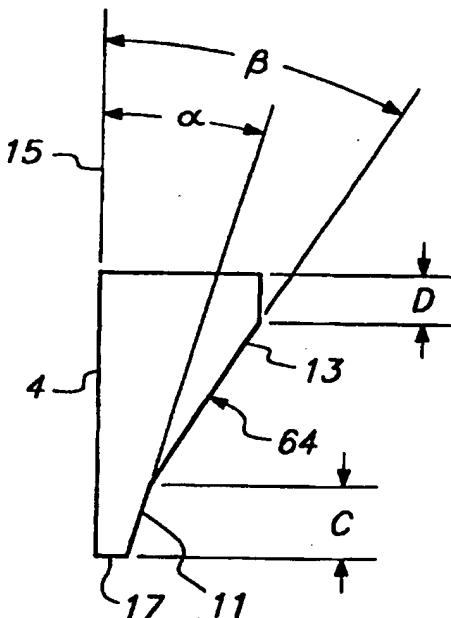
INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : A61N 1/30	A1	(11) International Publication Number: WO 97/48442 (43) International Publication Date: 24 December 1997 (24.12.97)
(21) International Application Number: PCT/US97/10596		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, AR IPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).
(22) International Filing Date: 18 June 1997 (18.06.97)		
(30) Priority Data: 60/019,990 18 June 1996 (18.06.96) US		
(71) Applicant: ALZA CORPORATION [US/US]; 950 Page Mill Road, P.O. Box 10950, Palo Alto, CA 94303-0802 (US).		
(72) Inventors: CORMIER, Michel, J., N.; 278 Andsbury Avenue, Mountain View, CA 94043 (US). NEUKERMANS, Armand, P.; 3510 Arbutus Avenue, Palo Alto, CA 94303 (US). BLOCK, Barry; 30610 Page Mill Road, Los Altos, CA 94022 (US). NAT, Avtar, S.; 34131 Siward Drive, Fremont, CA 94555-2128 (US).		Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(74) Agents: MILLER, Byron, D. et al.; Alza Corporation, 950 Page Mill Road, P.O. Box 10950, Palo Alto, CA 94303-0802 (US).		

(54) Title: DEVICE FOR ENHANCING TRANSDERMAL AGENT DELIVERY OR SAMPLING

(57) Abstract

A percutaneous agent delivery or sampling device (10) comprising a sheet (6) having at least one opening (8) therethrough and a plurality of microblades (4) for piercing the skin for increasing transdermal flux of an agent. The microblades (4) having a relatively sharp angled leading edge (11, 11') which transitions to a relatively gradually angled blade edge (13, 13').



FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

**DEVICE FOR ENHANCING TRANSDERMAL AGENT
DELIVERY OR SAMPLING**

TECHNICAL FIELD

6 The present invention relates to transdermal agent delivery and sampling.
7 More particularly, this invention relates to the transdermal delivery of agents,
8 such as peptides and proteins, as well as the transdermal sampling of agents, such as
9 glucose, body electrolytes and substances of abuse, such as but not limited to alcohol
10 and illicit drugs. The present invention uses skin-piercing microblades to enhance the
11 transdermal flux of the agents during transdermal delivery or sampling.

BACKGROUND ART

15 Interest in the percutaneous or transdermal delivery of peptides and proteins to
16 the human body continues to grow with the increasing number of medically useful
17 peptides and proteins becoming available in large quantities and pure form. The
18 transdermal delivery of peptides and proteins still faces significant problems. In many
19 instances, the rate of delivery or flux of polypeptides through the skin is insufficient to
20 produce a desired therapeutic effect due to the binding of the polypeptides to the skin.

21 In addition, polypeptides and proteins are easily degraded during and after
22 penetration into the skin, prior to reaching target cells. Likewise, the passive flux of
23 water soluble small molecules such as salts is limited.

24 One method of increasing the transdermal delivery of agents relies on the
25 application of an electric current across the body surface or on "electrotransport".
26 "Electrotransport" refers generally to the passage of a beneficial agent, e.g., a drug or
27 drug precursor, through a body surface such as skin, mucous membranes, nails, and
28 the like. The transport of the agent is induced or enhanced by the application of an
29 electrical potential, which results in the application of electric current, which delivers or
30 enhances delivery of the agent. The electrotransport of agents through a body

1 surface may be attained in various manners. One widely used electrotransport
2 process, iontophoresis, involves the electrically induced transport of charged ions.
3 Electroosmosis, another type of electrotransport process, involves the movement of a
4 solvent with the agent through a membrane under the influence of an electric field.
5 Electroporation, still another type of electrotransport, involves the passage of an agent
6 through pores formed by applying a high voltage electrical pulse to a membrane. In
7 many instances, more than one of these processes may be occurring simultaneously
8 to different extents. Electrotransport delivery generally increases agent delivery,
9 particularly large molecular weight species (e.g., polypeptides) delivery rates, relative
10 to passive or non-electrically assisted transdermal delivery. However, further
11 increases in transdermal delivery rates and reductions in polypeptide degradation
12 during transdermal delivery are highly desirable.

13 One method of increasing the agent transdermal delivery rate involves pre-
14 treating the skin with, or alternatively co-delivering with the beneficial agent, a skin
15 permeation enhancer. The term "permeation enhancer" is broadly used herein to
16 describe a substance which, when applied to a body surface through which the agent
17 is delivered, enhances its transdermal flux. The mechanism may involve an increase
18 in the permeability of the body surface, a reduction in the degradation of the agent
19 (e.g., degradation by skin enzymes) during transport, or in the case of electrotransport
20 delivery/sampling, a reduction of the electrical resistance of the body surface to the
21 passage of the agent therethrough or, the creation of hydrophilic pathways through the
22 body surface.

23 There have been many attempts to enhance transdermal flux by mechanically
24 puncturing the skin prior to transdermal drug delivery. See for example U.S. Patent
25 Nos. 5,279,544 issued to Gross et al., 5,250,023 issued to Lee et al., and 3,964,482
26 issued to Gerstel et al. These devices utilize tubular or cylindrical structures generally,
27 although Gerstel does disclose the use of other shapes, to pierce the outer layer of the
28 skin. Each of these devices provide manufacturing challenges, resistance to easy
29 penetration of the skin, and/or undesirable irritation of the skin.

1 As has been discussed, a variety of chemicals and mechanical means have
2 been explored to enhance transdermal flux. However, there is still a need to provide a
3 device suitable for increasing transdermal flux which device penetrates the skin with
4 very little insertion force, is low-cost and which can be manufactured reproducibly (i.e.,
5 without significant variation from device to device) in high volume production.

6

7 **DESCRIPTION OF THE INVENTION**

8

9 The present invention provides a reproducible, high volume production,
10 low-cost device capable of penetrating the skin easily and suitable for increasing
11 transdermal flux. The invention comprises a plurality of microblades for piercing the
12 skin having a leading edge with a relatively sharp angled first segment which
13 transitions to a relatively gradually angled second segment. The particular microblade
14 geometry allows better penetration of the skin with less "push down" (i.e., penetration
15 and insertion) force required of the user. The first segment forms a relatively small
16 angle with respect to an axis extending along the length of the microblade to provide a
17 very pointed section on the blade that pierces the skin readily. The leading edge then
18 transitions to a second segment which forms a larger angle relative to the axis than
19 the first segment. The second segment provides strength to the overall blade to
20 prevent bending due to the wider blade along that portion compared to the portion
21 along the first segment. The second segment, because of its larger width, also forms
22 longer slits in the skin thereby increasing the size of the transdermal pathways through
23 which agents can be delivered or withdrawn. Together, the sharper blade tip and the
24 relatively stronger blade base, improve the overall penetration characteristics of the
25 microblade and thereby reduce the push down force needed to achieve the desired
26 penetration quality.

27 The blades typically have a length of less than about 0.5 mm and a width and
28 thickness which is even smaller. In spite of their small size, the microblades can be
29 made with an extremely reproducible size and shape so that the microslits formed by

1 the microblades puncturing the skin also have a very reproducible size and depth.
2 Because the microblades have a small thickness (i.e., small relative to the width and
3 length of the blades), the microblades produce less tissue damage for a given
4 cross-section than a skin piercing microneedle having a circular cross-section. The
5 device of the present invention pierces the stratum corneum of a body surface to form
6 pathways through which a substance (e.g., a drug) can be introduced (i.e., delivery) or
7 through which a substance (e.g., a body electrolyte) can be withdrawn (i.e., sampling).

8 In one aspect of the invention, the device comprises a sheet having a plurality
9 of openings therethrough, a plurality of microblades integral therewith and extending
10 downward therefrom, at least a portion of the microblades having a leading edge with
11 a first angled segment and contiguous with the first angled segment a second angled
12 segment, the first angled segment being located distally on the microblade and having
13 a first angle relative to an axis along the length of the microblade, the second angled
14 segment having an angle greater relative to the axis than the first angle.

15 The device of the present invention can be used in connection with drug
16 delivery, body analyte or drug sampling, or both. Delivery devices for use with the
17 present invention include, but are not limited to, electrotransport devices, passive
18 devices, osmotic devices and pressure-driven devices. Sampling devices for use with
19 the present invention include, but are not limited to, "reverse" electrotransport devices
20 such as disclosed in Glikfeld et al., U.S. Patent No. 5,279,543 and Guy et al., U.S.
21 Patent No. 5,362,307, passive diffusion devices such as disclosed in Schoendorfer,
22 U.S. Patent No. 5,438,984, osmotic devices such as disclosed in Eckenhoff et al., U.S.
23 Patent No. 4,756,314, and negative pressure driven devices.

24

25 **BRIEF DESCRIPTION OF THE DRAWINGS**

26

27 Figure 1 is an enlarged perspective view of the skin proximal side of the
28 microblade array device in accordance with one embodiment of the present invention;

1 Figure 2 is an enlarged view of a portion of the microblades of the blade array
2 pattern;

3 Figure 3 is an enlarged view of a microblade in accordance with one
4 embodiment of the present invention;

5 Figures 4-7 are enlarged views of other embodiments of the microblade in
6 accordance with the present invention;

7 Figure 8 is a perspective exploded view of one embodiment of an
8 electrotransport agent delivery system with a microblade array device according to the
9 present invention; and

10 Figure 9 is a bottom plan view of the electrotransport agent delivery system of
11 figure 8.

12

13 **MODES FOR CARRYING OUT THE INVENTION**

14

15 Turning now to the drawings in detail, one embodiment of the skin-piercing
16 member 2 of the present invention is generally shown in Figure 1. Member 2 is used
17 in conjunction with percutaneous administration or sampling of an agent. The terms
18 "substance", "agent" and "drug" are used interchangeably herein and broadly include
19 physiologically or pharmacologically active substances for producing a localized or
20 systemic effect or effects in mammals including humans and primates, avians,
21 valuable domestic household, sport or farm animals, or for administering to laboratory
22 animals such as mice, rats, guinea pigs, and the like. These terms also include
23 substances such as glucose, body electrolytes, alcohol, illicit substances,
24 pharmaceuticals, illicit drugs, etc. that can be sampled through the skin. The major
25 barrier properties of the skin, such as resistance to drug penetration, reside with the
26 outermost layer (i.e., stratum corneum). The inner division of the epidermis generally
27 comprises three layers commonly identified as stratum granulosum, stratum malpighii,
28 and stratum germinativum. Once a drug penetrates below the stratum corneum, there
29 is substantially less resistance to permeation through the stratum granulosum, stratum

1 malpighii, and stratum germinativum. The device of the present invention is used to
2 form microslits in the stratum corneum and produce a percolation area in the skin for
3 improved transdermal delivery or sampling of an agent.

4 Member 2 comprises a plurality of microblades 4 (i.e., a blade array) extending
5 downward from one surface of a sheet or plate 6 (see Figure 1 in which a portion of
6 member 2 is in an inverted position to show the microblades). The microblades 4 are
7 sized and shaped to penetrate the stratum corneum of the epidermis when pressure is
8 applied to the device. The microblades form microslits in a body surface to increase
9 the administration of or sampling of a substance through the body surface. The term
10 "body surface" as used herein refers generally to the skin of an animal or human.

11 The microblades 4 are generally formed from a single piece of sheet material
12 and are sufficiently sharp and long for puncturing the stratum corneum of the skin. In
13 one embodiment, the microblades 4 and the sheet 6 are essentially impermeable or
14 are impermeable to the passage of an agent. The sheet 6 is formed with an opening 8
15 between the microblades 4 for enhancing the movement of an agent therethrough. In
16 the case of therapeutic agent (e.g., drug) delivery, the drug is released from a
17 drug-containing reservoir (not shown in Figure 2) through the opening 8 and passes
18 through microslits formed by the microblades 4 cutting through the stratum corneum,
19 migrates down the outer surfaces of the microblades and through the stratum corneum
20 to achieve local or systemic therapy. In the case of agent (e.g., body analyte)
21 sampling, the analyte migrates from the body through the microslits in the stratum
22 corneum which are cut by the microblades 4.

23 In one embodiment, the opening 8 corresponds to the portion of the sheet 6
24 occupied by each of the microblades prior to the blades being transpositioned into the
25 downward depending position. The number of microblades 4 per opening 8 can be
26 any number, preferably however between 1 and about 30 blades per opening.
27 Furthermore, the number of openings per device and the number of blades per device
28 are independent. The device may have only one opening and one microblade. The
29 agent can be administered at a controlled rate of release from the reservoir through an

1 agent release rate controlling material (not shown) covering the openings 8.

2 As is best shown in Figure 1, the microblades 4 have a thickness which is much
3 smaller than the width of the blades near their base, i.e., near the point where the
4 blades are attached to the sheet 6. This blade geometry provides maximum drug
5 percolation area with a minimum blade penetration area, and hence less tissue
6 damage. The drug percolation area is the skin area in contact with the blades which
7 provides for drug penetration in the skin. The microblades are shaped with the largest
8 possible surface area with a minimal cross-sectional area so as to give the largest
9 possible percolation area. Thin microblades are better than round protrusions for this
10 purpose because for the same cross-section, a thin blade produces more percolation
11 area and less tissue damage than a protrusion having a circular cross-section (i.e., a
12 cylindrically shaped protrusion). This is a crucial advantage over the prior art round
13 elements such as needles and tubes. Thin microblades also require less insertion
14 force than round protrusions. The width of each blade can be any of a range of
15 widths. The widths can be different from blade to blade in the array pattern. Likewise,
16 the width can be variable along the length of the blade, as will be described in more
17 detail below. The width of the blade at the intersection of the blade and the body
18 surface after the blade array has been inserted is preferably in the range of about 10
19 mm to about 500 mm, more preferably about 25 mm to about 400 mm, more
20 preferably 25 mm to about 300 mm.

21 The microblades 4 are provided with slanted (i.e., angled) leading edges 64
22 having multiple segments to reduce the insertion force required to press the blades
23 into the skin tissue. Because the blade insertion force is reduced, it is also possible to
24 use a thinner and more flexible sheet 6, which is advantageous in devices adapted to
25 be worn on the skin for extended (e.g. longer than 30 minutes) periods of time. In
26 Figures 1-5 and 7, the leading edges 64 have two segments each having a different
27 leading angle. The first segment 11 is the distal most segment. Contiguous with the
28 first segment 11 is second segment 13. The angle of the first segment relative to axis
29 or reference line 15 is designated as α . The angle of the second segment is

1 designated b. The multiple segmented slanted leading edge produces a cut through
2 the skin tissue that is equal to the full width of the blade 4 while reducing the amount
3 of metal that is in the skin tissue. In other words, a flat leading edge (i.e., a is 90°)
4 produces a blade with a larger amount of blade material in the skin tissue than is
5 produced by a blade having a slanted leading edge. The angle a of each segment 11
6 can be any angle between about 1° to 25°, preferably about 15°. The first segment 11
7 then transitions to the second segment 13 having an angle b between about 26° to
8 80°, preferably between about 30° to 45°, more preferably 35°.

9 The microblade 4 of the embodiments shown in Figures 4, 6 and 7 have sharp
10 distal tips 19 for easy penetration of the skin. The embodiments of microblade 4
11 shown in Figures 1-3 and 5 have a flattened distal most tip 17 which is easier to
12 manufacture and has greater resistance to bending upon insertion in the skin than the
13 more pointed tip 19.

14 The embodiments of Figures 1-4 and 6 have a single slanted leading edge 64
15 on the microblade 4, whereas the embodiments of Figures 5 and 7 have two slanted
16 leading edges 64 beginning approximately on the center line 15 and extending
17 outwardly therefrom on either side of the center line. As shown in Figure 7, the
18 slanted leading edges need not be symmetrical about the center line. First segment
19 11' is not equal to segment 11 and second segment 13' is not equal to segment 13.

20 The multiple segmented leading edge 64 of any of the embodiments previously
21 described can have any number of segments. For example, the embodiment of
22 Figure 6 has a third segment 21. The angle of the third segment 21 is designated g.
23 The second segment 13 transitions to the third segment 21 having an angle g relative
24 to reference line 15 greater than the angle b. Preferably, angle g is between about
25 35° to 80°, more preferably about 45°. As can be appreciated from Figure 6, a
26 plurality of contiguous angled segments wherein each of the subsequent angled
27 segments progressing proximally along the microblade from the first segment has an
28 angle relative to the reference line 15 greater than the angle of the preceding angled
29 segment creates a leading edge which appears arcuate (i.e., curved) in shape. In one

1 embodiment, the leading edge appears curved across the entire width of the blade.

2 The microblades 4 are formed using a photo-etching process. The photo-
3 etching process allows the microblades 4 to be reproducibly formed on a very small
4 (i.e., tens of microns) scale. This process also allows the microblades 4 to be formed
5 in shapes which require lower force for penetrating the skin. Some of the microblades
6 4 are provided with barbs 50 (Figures 1 and 2) in some fashion so that the member 2
7 and any corresponding device attached thereto stays attached to the skin after being
8 applied with pressure. The degree of attachment and the number and size of the
9 barbs 50 is such as to retain the delivery or sampling device during the normal activity
10 of the wearer, but not cause pain upon removal. As the microblades are pressed into
11 the skin tissue for use, the leading edge 64 of each microblade 4 cuts through and
12 pushes aside the skin tissue. After the microblades have come to rest in the skin, the
13 skin due to its elastic nature at least partially comes back together around the edges of
14 the microblades 4, in this way the surface 66 on each microblade having a barb 50
15 engages skin tissue and anchors the device in the skin. If the microblade is left in the
16 skin for an extended period of time (e.g., 24 hours), the skin tissue begins to heal
17 together in the area behind the surface 66 of the barb 50 thus improving the anchoring
18 of the device. Only one barb per blade is shown in the figures but it is within the scope
19 of the present invention that each blade can have a plurality of barbs extending
20 therefrom. The plurality of microblades 4 for puncturing the stratum corneum are
21 present on one face surface 48 of the member 2 in any predetermined arrangement,
22 for example, as a cluster of blades spaced in rows having any desired number, or in
23 any spaced apart relation of one blade to each other. Each blade has a width and
24 thickness that facilitates penetration of the stratum corneum without bending. In the
25 embodiment of Figure 1, there are six blades 4 along the perimeter of each opening 8
26 in sheet 6. Preferably, the width of each blade is between about 135 mm to about 300
27 mm and the length is about 600 mm. The required length of the blades is subject to
28 variation of the body surface being penetrated and corresponds to the natural
29 thickness of the stratum corneum, for one of the principle features of the invention is

1 that the blades are to penetrate the stratum corneum into the epidermis. Usually, the
2 blades will be about 25 mm to about 700 mm in length, with the length for most
3 applications being between about 50 mm to about 600 mm. By way of example, the
4 microblade 4 of Figure 3 is 254 mm wide and 508 mm in length wherein dimension C
5 is 127 mm and dimension D is 89 mm. The microblade 4 of Figure 4 is 254 mm wide
6 and 610 mm in length wherein dimension C is 127 mm and dimension D is 178 mm.
7 The sharp distal segment of the microblade is supported by the remainder of the blade
8 as it widens at an angle b and provides a relatively large base width which provides
9 the required structural integrity to prevent blade deflection upon insertion and
10 penetration in the skin.

11 The pattern for any of the blade array devices of the present invention is
12 produced with a photo-etching process. For example, reference may be had to U.S.
13 Provisional Application No. 60/019,990 filed June 18, 1996 of which any of the
14 disclosed methods can be used to produce the member 2 of the present invention. A
15 thin sheet or plate 6 of metal such as stainless steel or titanium is etched
16 photo-lithographically with patterns containing blade-like structures. In general, a thin
17 laminate dry resist or wet resist is applied on a sheet about 7 mm to about 100 mm
18 thick, preferably about 25 mm to about 50 mm thick. The resist is contact exposed
19 using a mask having the desired pattern and is subsequently developed. These
20 operations are conducted in much the same way that they are for the manufacture of a
21 printed circuit board. The sheet is then etched using acidic solutions. After the pattern
22 has been etched through the sheet, the sheet is placed on a die having a plurality of
23 openings corresponding to the openings 8 in the sheet. A punch having a plurality of
24 protrusions corresponding to the openings in the sheet and die is initially located
25 above the sheet and die. At the initial stage, the blades 4 are in the same plane as the
26 rest of the sheet 6. The protrusions on the punch are then pressed into the openings,
27 thus bending the blades 4 downward to be at an angle (e.g., substantially
28 perpendicular) to the plane of the sheet. The finished structure provides blades 4 with
29 an adjacent opening 8 for the passage of a substance therethrough when the member

1 2 is applied to the skin. Rectangular openings 8 are shown in the figures but the
2 invention encompasses the use of any shape openings including, but not limited to,
3 square, triangular, circular and elliptical. The blades 4 can be patterned with resist on
4 both sides of the sheet 6 and subsequently etched simultaneously from both sides to
5 achieve maximum pattern resolution for a given sheet thickness and to produce a
6 knife-like edge that can not be achieved with conventional stamping and punching
7 processes. Alternatively, the blades 4 can be patterned and etched from one side
8 only.

9 In another embodiment of the two-sided etching process, the blade array
10 pattern of any of the embodiments of the present invention is etched into the top
11 surface of sheet 6. A second pattern equivalent to the area bounded by each of the
12 openings 8 (e.g., rectangular) is etched into the bottom surface 48 such that each of
13 the blades in the blade array pattern is thinner than the surrounding sheet 6. As a
14 result, the sheet 6 forms a strong base and as the punch deforms the blades 4
15 downward, each of the blades plastically deforms so as to produce blades that are
16 straighter and more truly perpendicular to the sheet.

17 In one embodiment of the etching process, a dry resist (e.g., "Dynachem FL"
18 available from Dynachem located in Tustin, CA) is applied 12.5 mm thick to one or
19 both sides of the sheet and exposed in a standard manner. Then a suitable spray
20 etcher (e.g., "Dynamil VRP 1 0/NM" available from Western Tech. Assoc. located in
21 Anaheim, CA) is used to spray a mixture of ferric chloride and hydrochloric acid onto
22 the resist and sheet at 52 ° (125 °F) for two minutes. A standard caustic stripper is
23 used for the resist removal.

24 In another embodiment of the etching process, a wet resist (e.g., "Shipley
25 111S" available from Shipley Corporation, located in Marlborough, MA) is applied 7.5
26 mm thick at about 20 ° (70 °F) to one or both sides of the sheet and exposed in a
27 standard manner. Then a suitable etchant (e.g., ferric chloride) is sprayed onto the
28 resist and sheet at 49 ° (120 °F). A standard caustic stripper is used for the resist
29 removal.

1 Generally, the blades 4 are at an angle of about 90° to the surface 48 of the
2 sheet 6 after being punched, but they can be disposed at any angle forward or
3 backward from the perpendicular position that will facilitate penetration of and
4 attachment to the stratum corneum. In one embodiment, the blades are all aligned at
5 an angle between about 1° and about 89° degrees, preferably about 10° to about 60°,
6 more preferably about 20° to 45° to facilitate the device being slid along and into the
7 skin. The angled blades have two principal advantages. First, penetration of the
8 blades is not as strongly opposed by the elasticity of the skin because the blades are
9 slid generally horizontally into the skin as opposed to pressing vertically on the skin.
10 Second, the angled blades act to anchor the device in the skin as any motion of the
11 skin is less likely to dislodge the blades. In addition, other anchoring elements such as
12 barbs, openings, etc. can be used with the angled blades to further enhance
13 anchoring of the device.

14 The sheet and blades can be made from materials that have sufficient strength
15 and manufacturability to produce blades, such as, glasses, ceramics, rigid polymers,
16 metals and metal alloys. Examples of metals and metal alloys include but are not
17 limited to stainless steel, iron, steel, tin, zinc, copper, platinum, aluminum, germanium,
18 nickel, zirconium, titanium and titanium alloys consisting of nickel, molybdenum and
19 chromium, metals plated with nickel, gold, rhodium, iridium, titanium, platinum, and the
20 like. An example of glasses include a devitrified glass such as "PHOTOCERAM"
21 available from Corning in Corning, NY. Examples of polymers include but are not
22 limited to polystyrene, polymethylmethacrylate, polypropylene, polyethylene,
23 "BAKELITE", cellulose acetate, ethyl cellulose, styrene/acrylonitrile copolymers,
24 styrene/butadiene copolymers, acrylonitrile/butadiene/styrene (ABS) copolymers,
25 polyvinyl chloride and acrylic acid polymers including polyacrylates and
26 polymethacrylates.

27 The microblades of the present invention make an elongated, thin microcut
28 (i.e., a slit) in the skin surface because the blades have a small thickness (relative to
29 their width and length) resulting in a minimal blade cross-sectional area for the

1 portions of the blade in the skin. The geometry of the microblades 4 results in minimal
2 blade volume in the skin with maximal blade surface area in the skin. The advantages
3 of the present invention include, but are not limited to: (1) the very sharp first segments
4 on the leading edges make skin penetration easier; (2) the thin blade geometry
5 produces the maximum drug percolation area for a given cross-section of the blade;
6 (3) minimal tissue damage occurs because the amount of blade material in the skin
7 and hence the volume loading is minimized; (4) slanted leading edges (or equivalent
8 pointed shapes) further minimize the amount of volume loading or tissue damage
9 while preserving a large percolation area; (5) for a given volume loading, the larger the
10 surface area, the larger the frictional retaining force in the skin; and (6) for a given
11 desired percolation area, there are fewer blades necessary and therefore the force on
12 each tip is higher making skin penetration easier.

13 The number of blades and openings of any of the embodiments of the device 2
14 is variable with respect to the desired flux rate, agent being sampled or delivered,
15 delivery or sampling device used (i.e., electrotransport, passive, osmotic,
16 pressure-driven, etc.), and other factors as will be evident to one of ordinary skill in the
17 art. In general, the larger the number of blades per unit area (i.e., the blade density),
18 the more distributed is the flux of the agent through the skin because there are a
19 greater number of agent-conveying pathways through the skin. Consequently, the
20 smaller the number of blades per unit area, the more concentrated is the flux of the
21 agent through the skin because there are fewer pathways. The present invention has
22 a blade density of at least about 10 blades/cm² and less than about 1000 blades/cm²,
23 preferably at least about 600 blades/cm², more preferably at least about 800
24 blades/cm². In similar fashion, the number of openings per unit area through which
25 the agent passes is at least about 10 openings/cm² and less than about 1000
26 openings/cm². In one embodiment, the present invention produces a percolation area
27 of about 0.005 to .05 cm²/cm² of body surface, preferably about 0.01 cm²/cm² of body
28 surface.

1 One embodiment of the present invention relies on the application of an electric
2 current across the body surface or "electrotransport". Electrotransport refers generally
3 to the passage of a beneficial agent, e.g., a drug or drug precursor, through a body
4 surface such as skin, mucous membranes, nails, and the like. The transport of the
5 agent is induced or enhanced by the application of an electrical potential, which results
6 in the application of electric current, which delivers or enhances delivery of the agent
7 or, for "reverse" electrotransport, samples or enhances sampling of the agent. The
8 electrotransport of the agents into or out of the human body may be attained in various
9 manners. One widely used electrotransport process, iontophoresis, involves the
10 electrically induced transport of charged ions. Electroosmosis, another type of
11 electrotransport process involved in the transdermal transport of uncharged or
12 neutrally charged molecules (e.g., transdermal sampling of glucose), involves the
13 movement of a solvent with the agent through a membrane under the influence of an
14 electric field. Electroporation, still another type of electrotransport, involves the
15 passage of an agent through pores formed by applying an electrical pulse, a high
16 voltage pulse, to a membrane. In many instances, more than one of these processes
17 may be occurring simultaneously to different extents. Accordingly, the term
18 "electrotransport" is given herein its broadest possible interpretation, to include the
19 electrically induced or enhanced transport of at least one charged or uncharged agent,
20 or mixtures thereof, regardless of the specific mechanism(s) by which the agent is
21 actually being transported.

22 It will be appreciated by those working in the field that the present invention can
23 be used in conjunction with a wide variety of electrotransport systems, as the invention
24 is not limited in any way in this regard. For examples of electrotransport drug delivery
25 systems, reference may be had to U.S. Patent Nos. 5,147,296 to Theeuwes et al.,
26 5,080,646 to Theeuwes et al., 5,169,382 to Theeuwes et al., and 5,169,383 to Gyory
27 et al. For examples of "reverse" electrotransport devices, references may be had to
28 U.S. Patent Nos. 5,279,543 to Glikfeld et al. and 5,362,307 to Guy et al.

1 Electrotransport devices generally use at least two electrodes which are in
2 electrical contact with some portion of the skin, nails, mucous membranes, or other
3 body surface. In the case of transdermal agent delivery, one of the two electrodes is
4 commonly referred to as the "donor" or "active" electrode, and is the one from which
5 the agent is delivered into the body. In the case of transdermal agent sampling, one of
6 the two electrodes is referred to as the "receptor" electrode, and is the one into which
7 the agent (e.g., body analyte) is collected after being withdrawn from the body. The
8 second electrode is typically termed the "counter" or "return" electrode, and serves to
9 close the electrical circuit through the body. For example, when the agent to be
10 delivered is a cation, i.e., a positively charged ion, the anode becomes the active or
11 donor electrode, while the cathode serves to complete the circuit. Alternatively, if an
12 agent is an anion, i.e., a negatively charged ion, the cathode is the donor electrode.
13 When the agent to be sampled is a cation, the cathode becomes the receptor
14 electrode while the anode serves to complete the circuit. When the agent to be
15 sampled is an anion, the anode becomes the receptor electrode while the cathode
16 serves to complete the circuit. When the agent to be sampled has no net charge (e.g.,
17 glucose), then either the anode or the cathode, or both electrodes, can serve as the
18 receptor electrode. Both the anode and cathode may be donor electrodes if both
19 anionic and cationic agents are delivered simultaneously. Electrotransport delivery
20 systems generally require at least one reservoir or source of the agent to be delivered
21 to the body. Electrotransport sampling systems likewise require at least one reservoir
22 in which to collect the agent being sampled. Examples of such reservoirs include a
23 pouch or cavity as described in U.S. Patent No. 4,250,878 to Jacobsen, a porous
24 sponge or pad as described in U.S. Patent No. 4,141,359 to Jacobsen et al., and a
25 pre-formed gel body as described in U.S. Patent No. 4,383,529 to Webster, among
26 others. Such reservoirs are electrically connected to, and positioned between, the
27 anode or the cathode and the body surface, e.g. to provide a fixed or renewable
28 source of one or more drugs in the case of agent delivery. In addition, electrotransport
29 delivery systems also typically have an electrical power source, e.g., one or more

1 batteries, and an electrical controller designed to regulate the timing, amplitude and/or
2 frequency of the applied electric current, and hence regulate the timing and rate of
3 agent delivery/sampling. This power source component is electrically connected to
4 the two electrodes. Optional electrotransport device components include a counter
5 reservoir, adhesive coatings, insulating separation layers, and rate-controlling
6 membranes.

7 Figures 8 and 9 illustrate a representative electrotransport delivery/sampling
8 device 10 that may be used in conjunction with the present invention. Device 10
9 comprises an upper housing 16, a circuit board assembly 18, a lower housing 20,
10 anode electrode 22, cathode electrode 24, anode reservoir 26, cathode reservoir 28
11 and skin-compatible adhesive 30. Upper housing 16 has lateral wings 9 which assist
12 in holding device 10 on a patient's skin. Printed circuit board assembly 18 comprises
13 an integrated circuit 19 coupled to discrete components 40 and battery 32. Circuit
14 board assembly 18 is attached to housing 16 by posts (not shown in Figure 8) passing
15 through openings 13a and 13b, the ends of the posts being heated/melted in order to
16 heat stake the circuit board assembly 18 to the housing 16. Lower housing 20 is
17 attached to the upper housing 16 by means of adhesive layer 30, the upper surface 34
18 of adhesive layer 30 being adhered to both lower housing 20 and upper housing 16
19 including the bottom surfaces of wings 9. Shown (partially) on the underside of circuit
20 board assembly 18 is a button cell battery 32. Other types of batteries may also be
21 employed to power device 10 depending on the need.

22 The device 10 is generally comprised of battery 32, electronic circuitry 19,40,
23 electrodes 22,24, drug/receptor reservoir 26, counter reservoir 28, and member 2, all
24 of which are integrated into a self-contained unit. The outputs (not shown in Figure 8)
25 of the circuit board assembly 18 make electrical contact with the electrodes 24 and 22
26 through openings 23,23' in the depressions 25,25' formed in lower housing 20 by
27 means of electrically conductive adhesive strips 42,42'. Electrodes 22 and 24, in turn,
28 are in direct mechanical and electrical contact with the top sides 44',44 of drug
29 reservoirs 26 and 28. The bottom side 46 of drug reservoir 28 contacts the patient's

1 skin through the opening 29 in adhesive layer 30 (Figure 9). The bottom side 46' of
2 drug reservoir 26 contacts the patient's skin through the plurality of openings 8 in the
3 member 2. The formulation of reservoir 26 is preferably a viscous gel that fills the
4 openings 8 such that the reservoir 26 is in direct contact with the skin when the blades
5 have penetrated the stratum corneum. The contact between the reservoir and skin
6 provides a path for the agent to be transported along. If the reservoir 26 is not in direct
7 contact with the skin initially typically sweat accumulates in the confined area and
8 provides an agent-transmitting pathway between reservoir 26 and the skin.

9 Device 10 optionally has a feature which allows the patient to self-administer a
10 dose of drug, or self-sample a body electrolyte, by electrotransport. Upon depression
11 of push button switch 12, the electronic circuitry on circuit board assembly 18 delivers
12 a predetermined DC current to the electrode/reservoirs 22,26 and 24,28 for a delivery
13 interval of predetermined length. The push button switch 12 is conveniently located on
14 the top side of device 10 and is easily actuated through clothing. A double press of
15 the push button switch 12 within a short time period, e.g., three seconds, is preferably
16 used to activate the device, thereby minimizing the likelihood of inadvertent actuation
17 of the device 10. Preferably, the device transmits to the user a visual and/or audible
18 confirmation of the onset of operation by means of LED 14 becoming lit and/or an
19 audible sound signal from, e.g., a "beeper". Agent is delivered/sampled through the
20 patient's skin, e.g., on the arm, by electrotransport over the predetermined delivery
21 interval. Anodic electrode 22 is preferably comprised of silver and cathodic electrode
22 24 is preferably comprised of silver chloride. Both reservoirs 26 and 28 are preferably
23 comprised of polymeric gel materials. Electrodes 22,24 and reservoirs 26,28 are
24 retained by lower housing 20.

25 In the case of therapeutic agent (i.e., drug) delivery, a liquid drug solution or
26 suspension is contained in at least one of the reservoirs 26 and 28. Drug
27 concentrations in the range of approximately 1×10^{-4} M to 1.0 M or more can be used,
28 with drug concentrations in the lower portion of the range being preferred.

1 The push button switch 12, the electronic circuitry on circuit board assembly 18
2 and the battery 32 are adhesively "sealed" between upper housing 16 and lower
3 housing 20. Upper housing 16 is preferably composed of rubber or other elastomeric
4 material, e.g., injection moldable ethylene vinyl acetate. Lower housing 20 is
5 preferably composed of a plastic or elastomeric sheet material (e.g., polyethylene)
6 which can be easily molded to form depressions 25,25' and cut to form openings
7 23,23'. The assembled device 10 is preferably water resistant (i.e., splash proof) and
8 is most preferably waterproof. The system has a low profile that easily conforms to the
9 body, thereby allowing freedom of movement at, and around, the wearing site. The
10 reservoirs 26 and 28 are located on the skin-contacting side of the device 10 and are
11 sufficiently separated to prevent accidental electrical shorting during normal handling
12 and use.

13 The device 10 adheres to the patient's body surface (e.g., skin) by means of an
14 adhesive layer 30 (which has upper adhesive side 34 and body contacting adhesive
15 side 36). The adhesive side 36 covers the entire underneath side of the device 10
16 except where the member 2 and reservoir 28 are located. The adhesive side 36 has
17 adhesive properties which assures that the device 10 remains in place on the body
18 during normal user activity, and yet permits reasonable removal after the
19 predetermined (e.g., 24-hour) wear period. Upper adhesive side 34 adheres to lower
20 housing 20 and retains the electrodes and reservoirs within housing depression 25,25'
21 as well as retains member 2 to lower housing 20 and lower housing 20 to upper
22 housing 16.

23 In one embodiment of the drug delivery or sampling device there is a release
24 liner (not shown) on the device 10 for maintaining the integrity of the device when it is
25 not in use. In use, the release liner is stripped from the device before the device is
26 applied to the skin.

27 In other embodiments of the present invention, passive transdermal delivery or
28 sampling devices are used with member 2. In one embodiment the passive
29 transdermal delivery device comprises a reservoir containing agent. The reservoir is

1 preferably in the form of a matrix containing the agent dispersed therein. The reservoir
2 is sandwiched between a backing layer, which is preferably impermeable to the agent,
3 and a rate-controlling membrane. The reservoir is formed of a material, such as a
4 rubbery polymer, that is sufficiently viscous to maintain its shape. If a lower viscosity
5 material is used for the reservoir, such as an aqueous gel, the backing layer and
6 rate-controlling membrane would be sealed together about their periphery to prevent
7 leakage. In a sampling configuration, the reservoir would initially not contain the
8 agent. Located below the membrane is the microblade array member 2. The device
9 adheres to a body surface by means of a contact adhesive layer around the periphery
10 of the member 2. The adhesive layer may optionally contain agent. A strippable
11 release liner (not shown) is normally provided along the exposed surface of the
12 adhesive layer and is removed prior to application of the device to the body surface.

13 Alternatively, a transdermal therapeutic device in accordance with another
14 embodiment of the present invention can be attached to a body surface by means of a
15 flexible adhesive overlay. In this embodiment, the device is comprised of an
16 agent-containing reservoir (for a delivery configuration) which is preferably in the form
17 of a matrix containing the agent dispersed therein. In a sampling configuration, the
18 reservoir would initially not contain the agent. An impermeable backing layer is
19 provided adjacent one surface of the reservoir. The adhesive overlay maintains the
20 device on the body surface. The adhesive overlay can be fabricated together with, or
21 provided separately from, the remaining elements of the device. With certain
22 formulations, the adhesive overlay may be preferable to the contact adhesive
23 described previously. This is true, for example, where the agent reservoir contains a
24 material (such as, for example, an oily surfactant permeation enhancer) which
25 adversely affects the adhesive properties of the contact adhesive layer. The
26 impermeable backing layer is preferably slightly larger than the reservoir, and in this
27 manner prevents the agents in the reservoir from adversely interacting with the
28 adhesive in the overlay. Optionally, a rate-controlling membrane (not shown) can be
29 provided on the skin/mucosa side of the reservoir. A strippable release liner (not

1 shown) is also normally provided with the device and is removed just prior to
2 application of the device to the body surface.

3 The formulation for the passive transdermal devices may be aqueous or non-
4 aqueous based. The formulation is designed to deliver the drug at the necessary
5 fluxes. Aqueous formulations typically comprise water and about 1 to 2 weight percent
6 of a hydrophilic polymer as a gelling agent, such as hydroxyethylcellulose or
7 hydroxypropylcellulose. Typical non-aqueous gels are comprised of silicone fluid or
8 mineral oil. Mineral oil-based gels also typically contain 1 to 2 weight percent of a
9 gelling agent such as colloidal silicon dioxide.

10 The reservoir matrix should be compatible with the delivered agent, any
11 excipients (e.g., flux enhancers, irritation preventing agents) and/or any carrier
12 therefore. When using an aqueous-based system, the reservoir matrix is preferably a
13 hydrophilic polymer, e.g., a hydrogel. When using a non-aqueous-based system, the
14 reservoir matrix is preferably composed of a hydrophobic polymer. Suitable polymeric
15 matrices are well known in the transdermal drug delivery art.

16 The preferred form in which an agent is delivered or sampled generally
17 determines the type of delivery or sampling system to be used, and vice versa. That
18 is, the selection of a "passive" system which delivers or samples the agent by diffusion
19 or an electrically powered system which delivers or samples the agent by
20 electrotransport will be mostly determined by the form of the agent. For example, with
21 passive delivery systems, it has generally been recognized that the agent is preferably
22 delivered in either its free base or acid form, rather than in the form of a water soluble
23 salt. On the other hand, with electrotransport delivery devices, it has been recognized
24 that the drugs should preferably be ionized and the drug salt should be soluble in
25 water. For the case of pierced skin, there is substantial passive flux through the
26 microslits created by the microblades piercing the stratum corneum. For osmotic and
27 pressure driven systems which deliver or sample drugs by connective flow carried by a
28 solvent, the drug preferably has sufficient solubility in the carrier solvent. It will be
29 appreciated by those working in the field that the present invention can be used in

1 conjunction with a wide variety of osmotic delivery or sampling systems, as the
2 invention is not limited to a particular device in this regard. Osmotic devices are
3 disclosed for example in U.S. Patent Nos. 4,340,480 to Eckenhoff, 4,655,766 to
4 Theeuwes et al., and 4,753,651 to Eckenhoff.

5 This invention has utility in connection with the delivery of drugs within any of
6 the broad class of drugs normally delivered through body surfaces and membranes,
7 including skin. In general, this includes drugs in all of the major therapeutic areas.

8 The present invention has particular utility in the delivery of peptides,
9 polypeptides, proteins, nucleotidic drugs, and other such species through body
10 surfaces such as skin. These substances typically have a molecular weight of at least
11 about 300 daltons, and more typically have a molecular weight of at least about 300 to
12 40,000 daltons. Specific examples of peptides and proteins in this size range include,
13 without limitation, LHRH, LHRH analogs such as goserelin, buserelin, gonadorelin,
14 napharelin and leuprolide, GHRH, GHRF, insulin, insiltropin, calcitonin, octreotide,
15 endorphin, TRH, NT-36 (chemical name:
16 N-[(s)-4-oxo-2-azetidinyl]carbonyl]-L-histidyl-L-prolinamide), liprecin, pituitary
17 hormones (e.g., HGH, HMG, desmopressin acetate, etc.), follicle luteoids, aANF,
18 growth factors such as growth factor releasing factor (GFRF), bMSH, GH,
19 somatostatin, bradykinin, somatotropin, platelet-derived growth factor, asparaginase,
20 bleomycin sulfate, chymopapain, cholecystokinin, chorionic gonadotropin, corticotropin
21 (ACTH), erythropoietin, epoprostenol (platelet aggregation inhibitor), glucagon, HCG,
22 hirulog, hyaluronidase, interferon, interleukins, menotropins (urofollitropin (FSH) and
23 LH), oxytocin, streptokinase, tissue plasminogen activator, urokinase, vasopressin,
24 desmopressin, ACTH analogs, ANP, ANP clearance inhibitors, angiotensin II
25 antagonists, antidiuretic hormone agonists, bradykinin antagonists, ceredase, CSI's,
26 calcitonin gene related peptide (CGRP), enkephalins, FAB fragments, IgE peptide
27 suppressors, IGF-1, neurotrophic factors, colony stimulating factors, parathyroid
28 hormone and agonists, parathyroid hormone antagonists, prostaglandin antagonists,
29 pentigetide, protein C, protein S, renin inhibitors, thymosin alpha-1, thrombolytics,

1 TNF, vaccines, vasopressin antagonists analogs, alpha-1 antitrypsin (recombinant),
2 and TGF-beta.

3 As mentioned above, the member 2 of the present invention can also be used
4 with known sampling devices including, but not limited to, reverse iontophoresis,
5 osmosis, passive diffusion, phonophoresis, and suction (i.e., negative pressure).
6 Osmotic sampling devices can be used to sample any of a variety of agents through a
7 body surface including, but not limited to glucose, body electrolytes, alcohol, blood
8 gases, illicit drugs and illicit substances such as drugs of abuse. In another
9 embodiment, an osmotic sampling device is attached to a body surface by means of a
10 flexible adhesive overlay. The osmotic sampling device is comprised of a salt layer
11 located between a semi-permeable or osmotic membrane and an optional agent
12 sensing element. The optional agent sensing element can be any of a variety of
13 chemically reactive sensors and indicators, for example the color indicating test strips
14 associated with glucose testing. The adhesive overlay can have a cut-out or
15 transparent window in the area of the indicators so that the indicators can be readily
16 viewed. In an alternate embodiment, the agent sensing element can be located
17 between the osmotic sampling device and the salt layer.

18 While the invention has been described in conjunction with the preferred
19 specific embodiments thereof, it is to be understood that the foregoing description is
20 intended to illustrate and not limit the scope of the invention. Other aspects,
21 advantages and modifications within the scope of the invention will be apparent to
22 those skilled in the art to which the invention pertains.

1 **CLAIMS:**

2 1. A device (2) for piercing the stratum corneum of a body surface to form
3 pathways through which an agent can be introduced or withdrawn, comprising:

4 a sheet (6) having at least one opening (8) therethrough and a plurality of
5 microblades (4) extending downward therefrom, at least a portion of the plurality of
6 microblades (4) having a first angled segment (11,11') located distally on the
7 microblade (4) with a first angle (a) relative to an axis (15) along the length of the
8 microblade (4), and contiguous with the first angled segment (11,11'), a second angled
9 segment (13,13') having an angle (b) relative to the axis (15) greater than the first
10 angle a.

11
12 2. The device (2) of Claim 1, wherein the first angle (a) is in the range of about
13 1° to 25° and the angle (b) of the second angled segment (13,13') is in the range of
14 about 26° to 80°.

15
16 3. The device (2) of Claim 1, wherein the first angle (a) is about 15° and the
17 angle (b) of the second angled segment (13,13') is about 30°.

18
19 4. The device (2) of Claim 1, wherein the first angle (a) is about 15° and the
20 angle (b) of the second angled segment (13,13') is about 35°.

21
22 5. The device (2) of Claim 1, further comprising a third angled segment (21)
23 contiguous with the second angled segment (13,13') and having an angle (g) relative
24 to the axis (15) greater than the angle (b) of the second angled segment (13,13').

1 6. The device (2) of Claim 1, further comprising a plurality of contiguous angled
2 segments (11,11',13,13',21) wherein each of the subsequent angled segments
3 progressing proximally along the microblade (4) from the second angled segment
4 (13,13') have an angle (g) relative to the axis (15) greater than the angle (b) of the
5 preceding angled segment (13,13').

6

7 7. The device of claim 1, wherein the first and second angled segments
8 (11,13) are part of an arc-shaped microblade tip.

9

10 8. The device (2) of Claim 1, further comprising a therapeutic agent delivery
11 device (10) connected to the piercing device (2) and positioned to deliver a therapeutic
12 agent through the opening (8) to the body surface.

13

14 9. The device (2) of Claim 1, further comprising a sampling device (10)
15 connected to the piercing device (2) and positioned to sample a substance from the
16 body surface through the opening.

17

18 10. A device (2) for piercing the stratum corneum of a body surface to form
19 pathways through which an agent can be introduced or withdrawn, comprising:

20 a sheet (6) having at least one opening (8) therethrough and a plurality of
21 microblades (4) located along a periphery thereof and extending downward from the
22 sheet (6), at least a portion of the plurality of microblades (4) having first angled
23 segments (11,11') located distally on each side of an axis (15) along the length of the
24 microblade (4), each of the first segments (11,11') having a first angle (a) relative to
25 the axis (15), and contiguous with the first segments (11,11'), second segments
26 (13,13') on each side of the axis (15) each having an angle (b) relative to the axis (15)
27 greater than the first angle (a).

28

29

1 11. The device (2) of Claim 10, wherein the first angle (a) of each of the first
2 segments (11,11') is in the range of about 1° to 25° and the angle (b) of each of the
3 second angled segments (13,13') is in the range of about 26° to 80°.

4

5 12. The device (2) of Claim 10, wherein the first angle (a) of each of the first
6 segments (11,11') is about 15° and the angle (b) of each of the second segments
7 (13,13') is about 30°.

8

9 13. The device (2) of Claim 10, wherein the first angle (a) of each of the first
10 segments (11,11') are not equal angles.

11

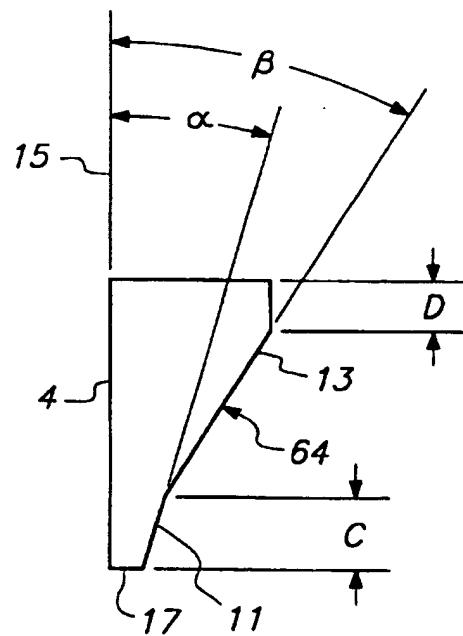
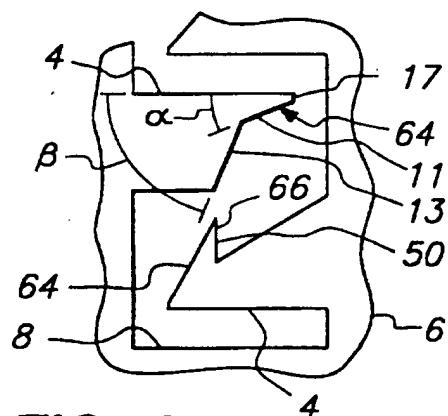
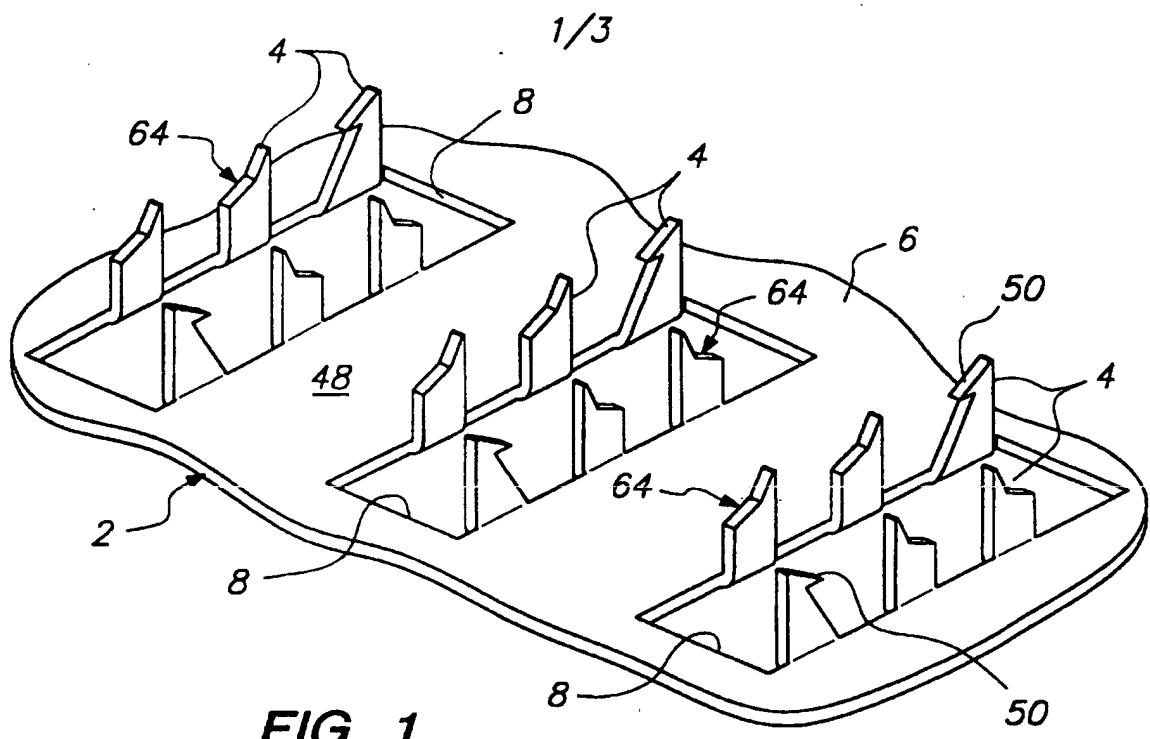
12 14. The device (2) of Claim 10, wherein the angle (b) of each of the second
13 segments (13,13') are not equal angles.

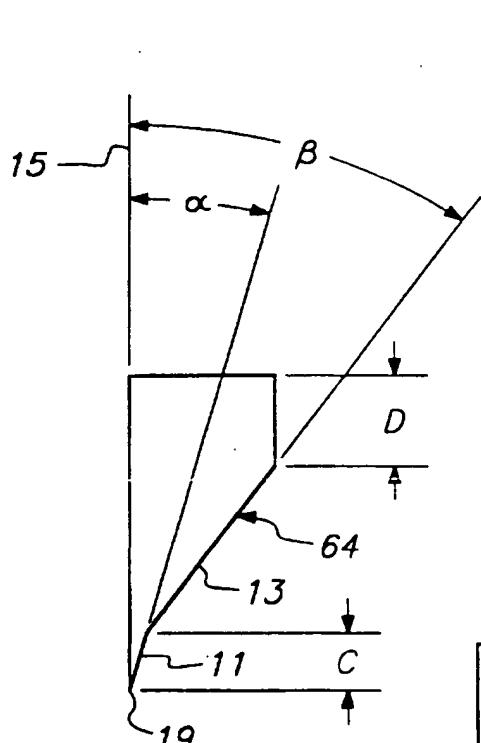
14

15 15. The device (2) of Claim 9, further comprising a therapeutic agent delivery
16 device (10) connected to the piercing device (2) and positioned to deliver a therapeutic
17 agent through the opening (8) to the body surface.

18

19 16. The device (2) of Claim 10, further comprising a sampling device (10)
20 connected to the piercing device (2) and positioned to sample a substance from the
21 body surface through the openings.





2/3

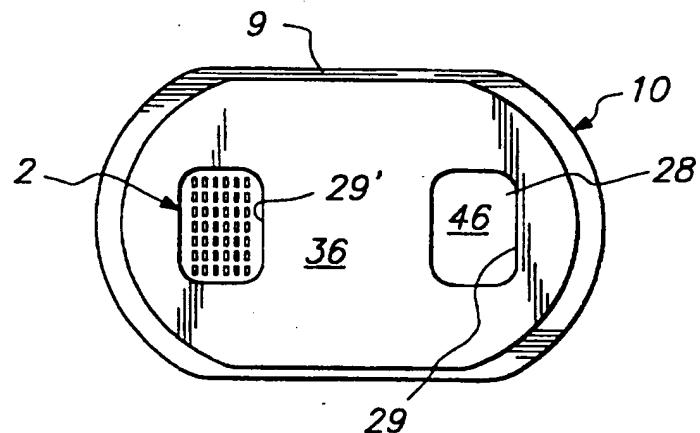
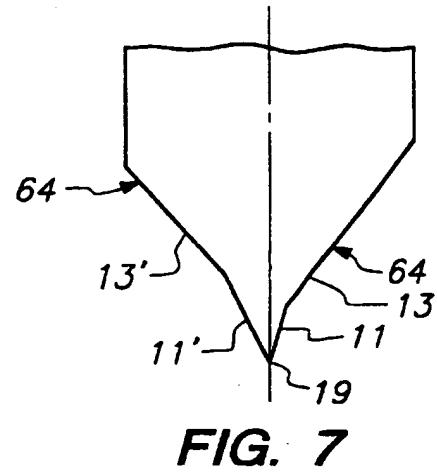
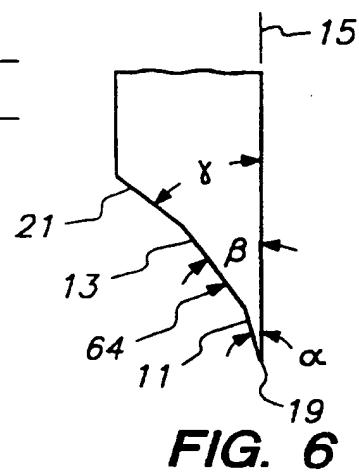
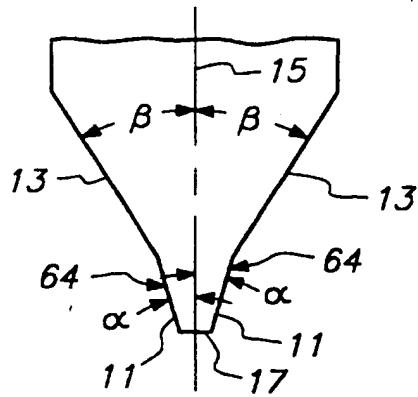
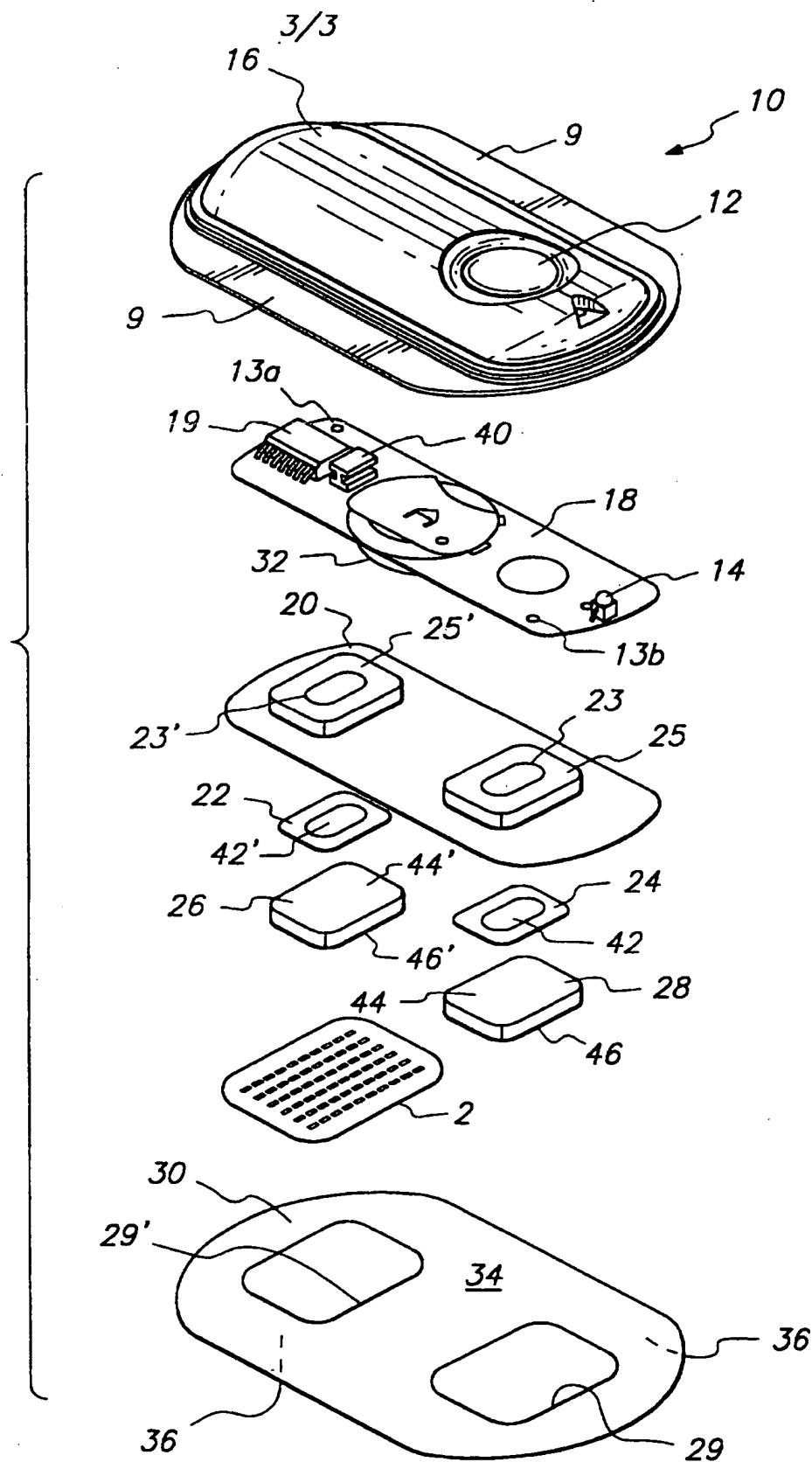


FIG. 8



INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 97/10596

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 A61N1/30

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 A61N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 96 17648 A (CIBA GEIGY AG ;EFFENHAUSER CARLO STEFAN (DE); MANZ ANDREAS (CH)) 13 June 1996	1-4,8, 10-12
Y	see page 7, line 28 - page 12, line 27; figures ---	9
Y	US 5 279 543 A (GLIKFELD ET AL.) 18 January 1994 cited in the application	9
A	see column 3, line 64 - column 4, line 49; figures 7,8 ---	1,10,16
A	US 5 279 544 A (GROSS ET AL.) 18 January 1994 cited in the application see page 2, line 58 - page 7, line 10; figures ---	1-4,8, 10-12,15
		-/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

& document member of the same patent family

1

Date of the actual completion of the international search

22 October 1997

Date of mailing of the international search report

10. 11. 97

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl
Fax: (+31-70) 340-3016

Authorized officer

Rakotondrajaona, C

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 97/10596

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 3 964 482 A (GERSTEL MARTIN S ET AL) 22 June 1976 cited in the application see column 4, line 21 - column 7, line 20; figures ---	1,8,10, 15
A	US 5 250 023 A (LEE ET AL.) 5 October 1993 cited in the application see column 1, line 29 - column 11, line 8; figures ---	1,8,10, 15
A	WO 92 10234 A (LIFSCHITZ) 25 June 1992 see page 4, line 10 - page 7, line 25; figures -----	1,8,10

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/US 97/10596

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9617648 A	13-06-96	AU 4256496 A EP 0796128 A	26-06-96 24-09-97
US 5279543 A	18-01-94	AT 129909 T AU 639888 B AU 3183889 A DE 68924716 D DE 68924716 T DK 179390 A EP 0326398 A EP 0673622 A ES 2085863 T IE 63406 B JP 4502561 T KR 9711449 B PT 89560 B WO 8906989 A US 5362307 A	15-11-95 12-08-93 25-08-89 14-12-95 04-07-96 28-07-90 02-08-89 27-09-95 16-06-96 19-04-95 14-05-92 11-07-97 31-05-95 10-08-89 08-11-94
US 5279544 A	18-01-94	US 5156591 A AT 156375 T DE 69312916 D WO 9317754 A EP 0630276 A JP 7508427 T US 5527288 A AT 142115 T AU 642112 B AU 9058791 A DE 69121881 D DE 69121881 T WO 9210234 A EP 0516783 A JP 5504711 T NZ 240875 A ZA 9301775 A	20-10-92 15-08-97 11-09-97 16-09-93 28-12-94 21-09-95 18-06-96 15-09-96 07-10-93 08-07-92 10-10-96 03-04-97 25-06-92 09-12-92 22-07-93 27-04-94 30-09-93
US 3964482 A	22-06-76	NONE	
US 5250023 A	05-10-93	EP 0429842 A	05-06-91

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/US 97/10596

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5250023 A		EP 0509122 A JP 1892430 C JP 3151982 A JP 6014980 B CA 2041250 A,C	21-10-92 26-12-94 28-06-91 02-03-94 23-11-91
WO 9210234 A	25-06-92	US 5156591 A AT 142115 T AU 642112 B AU 9058791 A DE 69121881 D DE 69121881 T EP 0516783 A JP 5504711 T NZ 240875 A US 5527288 A US 5279544 A	20-10-92 15-09-96 07-10-93 08-07-92 10-10-96 03-04-97 09-12-92 22-07-93 27-04-94 18-06-96 18-01-94